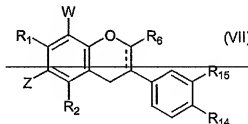
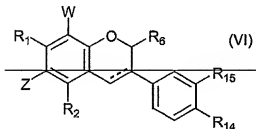
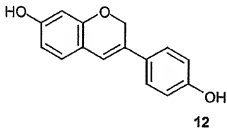


**AMENDMENTS TO THE CLAIMS**

**This listing of claims will replace all prior versions and listings of claims in the application:**

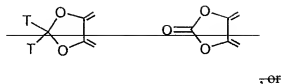
**LISTING OF CLAIMS:**

1. (currently amended): A method of increasing the sensitivity of cancer cells or a tumour to a chemotherapeutic agent by contacting said cells or tumour with an isoflavonoid compound of formula 12(VI) or (VII):

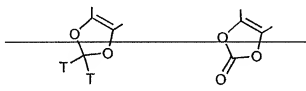


wherein

R<sub>1</sub>, R<sub>2</sub> and Z are independently hydrogen, hydroxy, OR<sub>9</sub>, OC(O)R<sub>10</sub>, OS(O)R<sub>10</sub>, CHO, C(O)R<sub>10</sub>, COOH, CO<sub>2</sub>R<sub>10</sub>, CONR<sub>3</sub>R<sub>4</sub>, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl, alkylaryl, alkoxyaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R<sub>2</sub> is as previously defined, and R<sub>1</sub> and Z taken together with the carbon atoms to which they are attached form a five-membered ring selected from



$R_1$  is as previously defined, and  $R_2$  and  $Z$  taken together with the carbon atoms to which they are attached form a five-membered ring selected from



$W$  is  $R_4$ ;

$R_3$  is hydrogen, alkyl, arylalkyl, alkenyl, aryl, an amino acid,  $C(O)R_{11}$  where  $R_{11}$  is hydrogen, alkyl, aryl, arylalkyl or an amino acid, or  $CO_2R_{12}$  where  $R_{12}$  is hydrogen, alkyl, haloalkyl, aryl or arylalkyl;

$R_4$  is hydrogen, alkyl or aryl, or

$R_3$  and  $R_4$  taken together with the nitrogen to which they are attached comprise pyrrolidinyl or piperidinyl;

$R_6$  is hydrogen, hydroxy, alkyl, aryl, amino, thio,  $NR_3R_4$ ,  $COR_{11}$  where  $R_{11}$  is as previously defined,  $CO_2R_{12}$  where  $R_{12}$  is as previously defined or  $CONR_3R_4$ ;

$R_9$  is alkyl, haloalkyl, aryl, arylalkyl,  $C(O)R_{11}$  where  $R_{11}$  is as previously defined, or  $Si(R_{13})_3$  where  $R_{13}$  where each  $R_{13}$  is independently hydrogen, alkyl or aryl;

$R_{10}$  is hydrogen, alkyl, haloalkyl, amino, aryl, arylalkyl, an amino acid, alkylamino or dialkylamino;

the drawing “” represents either a single bond or a double bond;

~~T is independently hydrogen, alkyl or aryl,~~

~~R<sub>14</sub> and R<sub>15</sub> are independently hydrogen, hydroxy, OR<sub>9</sub>, OC(O)R<sub>10</sub>, OS(O)R<sub>10</sub>, CHO, C(O)R<sub>10</sub>,~~

~~COOH, CO<sub>2</sub>R<sub>10</sub>, CONR<sub>3</sub>R<sub>4</sub>, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl,~~

~~thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R<sub>14</sub> and R<sub>15</sub> are fused~~

~~together to form a cyclic alkyl, aromatic or heteroaromatic structure,~~

~~and pharmaceutically acceptable salts thereof, and,~~

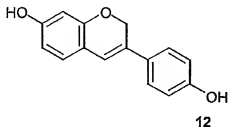
~~wherein the cancer is ovarian, pancreatic or prostate cancer, and~~

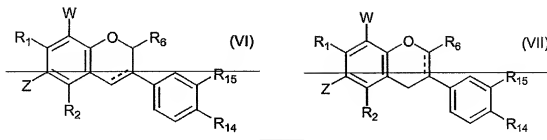
~~the chemotherapeutic agent is platinum-based or anti-mitotic agent cisplatin, carboplatin,~~  
~~paclitaxel, gemcitabine or doxorubicin.~~

2. (previously presented): A method of claim 1, wherein prior to the contacting, the cancer cells or tumour were/was not sensitive to the chemotherapeutic agent.

3. (currently amended): A method of claim 1, wherein the compound of formula 12(VI) or (VII) is administered to a subject in need of such treatment.

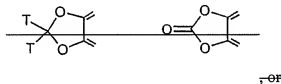
4. (currently amended): A combination therapy for the treatment or prophylaxis of ~~cell proliferation, cancer or a disease associated with oxidant stress~~ comprising administering to a subject a therapeutically effective amount of a compound of formula 12(VI) or (VII) and a chemotherapeutic agent:



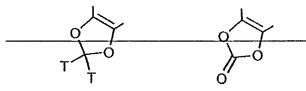


wherein

R<sub>1</sub>, R<sub>2</sub> and Z are independently hydrogen, hydroxy, OR<sub>3</sub>, OC(O)R<sub>10</sub>, OS(O)R<sub>10</sub>, CHO, C(O)R<sub>10</sub>, COOH, CO<sub>2</sub>R<sub>10</sub>, CONR<sub>3</sub>R<sub>4</sub>, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl, alkylaryl, alkoxyaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R<sub>2</sub> is as previously defined, and R<sub>1</sub> and Z taken together with the carbon atoms to which they are attached form a five-membered ring selected from



R<sub>1</sub> is as previously defined, and R<sub>2</sub> and Z taken together with the carbon atoms to which they are attached form a five-membered ring selected from



W is R<sub>1</sub>,

~~R<sub>3</sub> is hydrogen, alkyl, arylalkyl, alkenyl, aryl, an amino acid, C(O)R<sub>11</sub> where R<sub>11</sub> is hydrogen, alkyl, aryl, arylalkyl or an amino acid, or CO<sub>2</sub>R<sub>12</sub> where R<sub>12</sub> is hydrogen, alkyl, haloalkyl, aryl or arylalkyl,~~

~~R<sub>4</sub> is hydrogen, alkyl or aryl, or~~

~~R<sub>3</sub> and R<sub>4</sub> taken together with the nitrogen to which they are attached comprise pyrrolidinyl or piperidinyl,~~

~~R<sub>6</sub> is hydrogen, hydroxy, alkyl, aryl, amino, thio, NR<sub>3</sub>R<sub>4</sub>, COR<sub>11</sub> where R<sub>11</sub> is as previously defined, CO<sub>2</sub>R<sub>12</sub> where R<sub>12</sub> is as previously defined or CONR<sub>3</sub>R<sub>4</sub>,~~

~~R<sub>9</sub> is alkyl, haloalkyl, aryl, arylalkyl, C(O)R<sub>11</sub> where R<sub>11</sub> is as previously defined, or Si(R<sub>13</sub>)<sub>3</sub> where R<sub>13</sub> where each R<sub>13</sub> is independently hydrogen, alkyl or aryl,~~

~~R<sub>10</sub> is hydrogen, alkyl, haloalkyl, amino, aryl, arylalkyl, an amino acid, alkylamino or dialkylamino,~~

~~the drawing "—" represents either a single bond or a double bond,~~

~~T is independently hydrogen, alkyl or aryl,~~

~~R<sub>14</sub> and R<sub>15</sub> are independently hydrogen, hydroxy, OR<sub>9</sub>, OC(O)R<sub>10</sub>, OS(O)R<sub>10</sub>, CHO, C(O)R<sub>10</sub>, COOH, CO<sub>2</sub>R<sub>10</sub>, CONR<sub>3</sub>R<sub>4</sub>, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R<sub>14</sub> and R<sub>15</sub> are fused together to form a cyclic alkyl, aromatic or heteroaromatic structure,~~

~~and pharmaceutically acceptable salts thereof, and,~~

~~wherein the cancer is ovarian, pancreatic or prostate cancer, and~~

~~the chemotherapeutic agent is platinum-based or anti-mitotic agent cisplatin, carboplatin, paclitaxel, gemcitabine or doxorubicin.~~

5.-7. (canceled).

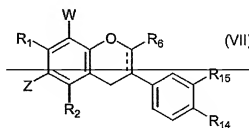
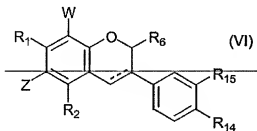
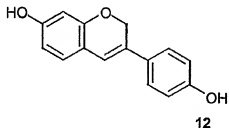
8. (currently amended): A method of claim 4, wherein the administration of the compound of formula 12(VI) or (VII) precedes the administration of the chemotherapeutic agent.

9. (currently amended): A method of claim 4, wherein the administration of the compound of formula 12(VI) or (VII) and the chemotherapeutic agent is simultaneous.

10. (currently amended): A method claim 4, wherein the combination therapy follows observed resistance by cancer cells or tumour to [[a]]the chemotherapeutic agent.

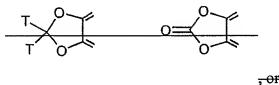
11.-22. (canceled).

23. (currently amended): A pharmaceutical composition comprising a compound of formula 12(VI) or (VII) and a chemotherapeutic agent:

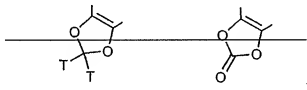


wherein

$R_1$ ,  $R_2$  and  $Z$  are independently hydrogen, hydroxy, OR, OC(O) $R_{10}$ , OS(O) $R_{10}$ , CHO, C(O) $R_{10}$ , COOH, CO<sub>2</sub> $R_{10}$ , CONR<sub>3</sub> $R_4$ , alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl, alkylaryl, alkoxyaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or  $R_2$  is as previously defined, and  $R_1$  and  $Z$  taken together with the carbon atoms to which they are attached form a five-membered ring selected from



$R_1$  is as previously defined, and  $R_2$  and  $Z$  taken together with the carbon atoms to which they are attached form a five-membered ring selected from



$W$  is  $R_1$ ,

$R_3$  is hydrogen, alkyl, arylalkyl, alkenyl, aryl, an amino acid, C(O) $R_{11}$ , where  $R_{11}$  is hydrogen, alkyl, aryl, arylalkyl or an amino acid, or CO<sub>2</sub> $R_{12}$ , where  $R_{12}$  is hydrogen, alkyl, haloalkyl, aryl or arylalkyl,

$R_4$  is hydrogen, alkyl or aryl, or

$R_3$  and  $R_4$  taken together with the nitrogen to which they are attached comprise pyrrolidinyl or piperidinyl,

$R_6$  is hydrogen, hydroxy, alkyl, aryl, amino, thio, NR<sub>3</sub> $R_4$ , COR<sub>11</sub>, where  $R_{11}$  is as previously defined, CO<sub>2</sub> $R_{12}$ , where  $R_{12}$  is as previously defined or CONR<sub>3</sub> $R_4$ ,

~~R<sub>9</sub> is alkyl, haloalkyl, aryl, arylalkyl, C(O)R<sub>11</sub>, where R<sub>11</sub> is as previously defined, or Si(R<sub>13</sub>)<sub>3</sub>~~

~~where R<sub>13</sub>, where each R<sub>13</sub> is independently hydrogen, alkyl or aryl,~~

~~R<sub>10</sub> is hydrogen, alkyl, haloalkyl, amino, aryl, arylalkyl, an amino acid, alkylamino or dialkylamino,~~

~~the drawing "—" represents either a single bond or a double bond,~~

~~T is independently hydrogen, alkyl or aryl,~~

~~R<sub>14</sub> and R<sub>15</sub> are independently hydrogen, hydroxy, OR<sub>9</sub>, OC(O)R<sub>10</sub>, OS(O)R<sub>10</sub>, CHO, C(O)R<sub>10</sub>,~~

~~COOH, CO<sub>2</sub>R<sub>10</sub>, CONR<sub>3</sub>R<sub>4</sub>, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl,~~

~~thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R<sub>14</sub> and R<sub>15</sub> are fused together to form a cyclic alkyl, aromatic or heteroaromatic structure,~~

~~and pharmaceutically acceptable salts thereof, and~~

~~wherein the chemotherapeutic agent is platinum-based or anti-mitotic agent cisplatin, carboplatin, paclitaxel, gemcitabine or doxorubicins.~~

24.-28. (Canceled).